

The Upper Functional G.I. Disorder

# The Pseudo-ulcer



Ulcer-like symptoms: no G.I. pathology

The patient is convinced it's an ulcer. However, symptoms are not quite typical, and x-ray findings are negative. These findings and the results of additional diagnostic procedures exclude an organic basis for the patient's complaints. A diagnosis of "upper functional gastrointestinal disorder" is made, which is supported by the fact that episodes of painful symptoms coincide with episodes of excessive anxiety, as indicated by the history.

It may be useful to explain to the patient the mechanism by which emotions upset normal G.I. functioning, resulting in hypersecretion and hypermotility and thus causing such symptoms as nausea and epigastric pain. In upper functional gastrointestinal disorders, counseling by the primary physician can often help the patient to understand and how excessive anxiety may cause flare-ups of G.I. symptoms.

A disproportionate number of patients seen by the general practitioner suffer from functional disorders, as do more than half of those seen by the gastroenterologist.\* Where milder cases may respond to counsel-

ing alone, if symptoms are severe and disabling to any degree, a suitable regimen may include medication to reduce the symptoms and the excessive anxiety that often provokes these distressing symptoms. In these cases, Librax as an adjunct can greatly contribute to the course of therapy. Its dual action can offer relief of both painful symptoms and excessive anxiety, because each capsule contains 5 mg chloridazepoxide HCl and 2.5 mg cimetidine Br. The antianxiety

An adjunct  
in anxiety-related upper  
functional G.I. disorders

## Librax®

Each capsule contains 5 mg chloridazepoxide HCl and 2.5 mg cimetidine Br.

among drugs for certain gastrointestinal disorders associated with excessive anxiety: the cimetidine bromide (Quarzan™) component furnishes dependable antisecretory-antispasmodic action. Dosage is flexible; it may be adjusted according to your patient's requirements within the range of 1 or 2 capsules three or four times daily, up to 8 capsules daily in divided doses.

\*Rome HP, Brannick TL: Orientation and mechanism of functional disorders: clinicalphysiologic correlation, chap. 18, in *Gastroenterology*, edited by Beckus HL, Philadelphia, WB Saunders Company, 1965, p. 1116

pregnancy, lactation, or in women of childbearing age requires that its potential benefits be weighed against its possible hazards. As with all anticholinergic drugs, an inhibiting effect on lactation may occur.

Precautions: In elderly and debilitated, limit dosage to smallest effective amount to preclude development of ataxia, dizziness or confusion (not more than two capsules per day initially; increase gradually as needed and tolerated). Though generally not recommended, if combination therapy with other psychotropics seems indicated, carefully consider individual pharmacologic effects, particularly in use of potent tranquilizing drugs such as MAO inhibitors and phenothiazines. Observe usual precautions in presence of impaired renal or hepatic function. Paradoxical reactions (e.g., excitement, stimulation) and acute rage have been reported in psychiatric patients. Therapy used cautiously in treatment of anxiety states with evidence of impending depression; suicidal tendency may be present and protective measures necessary. Variable effects on blood coagulation have been reported; very rarely, relationship has not been established clinically.

Adverse Reactions: No side effects or manifestations not seen with either compound alone have been reported with Librax. When chloridazepoxide hydrochloride is used alone, drowsi-

ness, ataxia and confusion may occur, especially in the elderly. These are reversible in most instances by proper dosage adjustment, but are also occasionally observed at the lower dosage ranges. In a few instances symptoms have been reported. Also encountered are isolated instances of skin eruptions, edema, minor menstrual irregularities, nausea and constipation, extrapyramidal symptoms, increased and decreased libido—all infrequent and generally controlled with dosage reduction; changes in EEG patterns (low-voltage dysrhythmia including agranulocytosis), jaundice and hepatic dysfunction have been reported occasionally with chloridazepoxide hydrochloride, making periodic blood counts and liver function tests advisable during protracted therapy. Adverse effects reported with Librax are typical of anticholinergic agents, i.e., dryness of mouth, blurring of vision, urinary hesitancy and constipation. Constipation has occurred most often when Librax therapy is combined with other "spasmolytics" and/or low residue diets.

Roche Laboratories  
Division of Hoffmann-La Roche Inc.  
Nutley, New Jersey 07110



Before prescribing, please consult complete product information, a summary of which follows:

Indications: Symptomatic relief of hypersecretion, hypermotility and anxiety and tension states associated with organic or functional gastrointestinal disorders and as adjunctive therapy in the management of peptic ulcer, gastritis, duodenitis, irritable bowel syndrome, spastic colitis, and mild ulcerative colitis.

Contraindications: Patients with glaucoma; prostatic hypertrophy and benign bladder neck obstruction; known hypersensitivity to chloridazepoxide hydrochloride and/or cimetidine bromide.

Warnings: Caution patients about possible combined effects with alcohol and other CNS depressants. As with all CNS-acting drugs, caution patients against hazardous occupational requirements, complete mental alertness (e.g., operating machinery, driving). Though physical and psychological dependence have rarely been reported on recommended doses, use caution in administering Librax (chloridazepoxide hydrochloride) to known addiction-prone individuals or those who might develop addiction. Withdrawal symptoms (including convulsions, increased dosage, withdrawal symptoms) following convulsions, following discontinuation of the drug and similar to those seen with barbiturates, have been reported. Use of any drug in

A B C D

# Medical Tribune

and Medical News

with news of medicine and its practice by separate complete

Behind News: CIR Delegates Discussing Strike Talks



Dr. Jay Debbin (standing at right in T-shirt), chairman of negotiating committee of Committee of Interns and Residents, reports to C.I.R. delegate caucus on progress of talks during strike affecting 23 New York City hospitals.

## New York Strike May Set Pattern For Hospital Staff Work Changes

Medical Tribune Report

NEW YORK—Last month's strike by some 2,100 interns and residents in New York City hospitals didn't last long, but it was watched closely around the country and the terms of the settlement that was reached may serve as precedent for changes in other cities.

Although the strike was a success from the point of view of the young doctors who led and participated in it, interviews conducted by MEDICAL TRIBUNE suggest that strikers and non-strikers alike had mixed feelings about the action.

The strike was called by the Committee of Interns and Residents, representing health officers at city and voluntary health care facilities in the greater New York area, against the League of Voluntary Hospitals, a

group of 11 hospital centers and their municipal affiliates. At issue were house staff duty schedules—as much as 58 hours a week, with weekly on-duty time often exceeding 100 hours for junior house staff.

Another area of disagreement was so-called "out-of-title" work. In many cases, it was alleged by the C.I.R., interns and residents were forced to do the work of nurses, technicians, aides, and even messengers because of understaffing.

According to the terms of the settlement, no later than July 1, 1976, no house staff officer will be required to perform call duty more than one night in three. Also, by May 1, 1975, standing committees will be set up in each

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## NIH Study Finds Nitroglycerin Beneficial in Acute Infarction

By HARRIET PAGE  
Medical Tribune Staff

BETHESDA, Mo.—The use of nitroglycerin is proving to be "consistently beneficial" in the treatment of patients with acute myocardial infarction, according to Dr. Stephen E. Epstein, of the Cardiology Branch of the National Heart and Lung Institute.

Twelve patients have so far been treated in a collaborative study headed by Dr. Epstein. And so far, he told MEDICAL TRIBUNE, with a follow-up of up to six months, the clinical response is bearing out the beneficial results he and his associates found in earlier animal studies.

"Nitroglycerin appears to reverse the manifestations of heart failure, reduce the size of the infarct, and permit the heart to pump more effectively," Dr. Epstein said. But he warned that these are still preliminary results and that the administration of nitroglycerin must be carefully monitored.

He and his associates have been giving what he termed multiple "large" doses sublingually over a 15-minute period, Dr. Epstein said, but at the first

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## Briton Fears Pool Of Hepatitis B Virus Rising in Newborn

By FRANCES GODONWORTH  
Medical Tribune Staff

NEW YORK—A pool of "chronic carriers" of hepatitis B virus may be building up among children born to women who became infected with it during pregnancy or who are asymptomatic carriers themselves, a British investigator warned here.

Calling the situation a "cause of utmost concern," Dr. Aric J. Zuckerman, of the London School of Hygiene and Tropical Medicine, urged that all pregnant women be screened routinely for hepatitis B surface antigen just as blood donors are tested.

The virologist noted that one in 1,000 healthy volunteer blood donors in Britain are carriers of hepatitis B antigen—one in 500 among certain social groups—and that the carrier incidence reaches 20 per cent in some countries.

The finding of active infection or a carrier state in pregnant women should be the signal to undertake measures to protect the child, Dr. Zuckerman told a symposium on infections of the fetus and newborn presented by the New York University School of Medicine

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## 1969-73 Study Results

## Control Program Reduces Hospital Infections by 10%

Medical Tribune Report

ROCKFORD, ILL.—What is believed to be the first long-term study to gauge the overall effect of hospital infection control on morbidity and mortality, and to show that the rate of such infections can be reduced, was described by Dr. Larry D. Edwards, Associate Professor of Medicine and chief of infectious diseases at Rockford School of Medicine, in a recent interview.

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making rounds at press time

SELL YOUR KIDNEY? An unemployed Pittsburg man deeply in debt tried it and had a prospective buyer, until Dr. Keith Hruska, co-director of the Chronology American Kidney Center, Washington, U.S. St. Louis, talked them out of it because it probably was not a suitable match. "I can

understand the seller's desperation — and there appears to be an increase lately in attempts to sell organs — but we have to avoid a situation in which the rich can buy their health and the poor cannot," Dr. Hruska told MT. He attributed the increase in sales attempts to the ailing economy. Dr. Ira Grier, medical director of the National Kidney Foundation added that "buying and selling organs will create a black market, with sales to the highest bidder."

OPPOSED — H.E.W. is "paying close attention," a spokesman told MT to some 2,300 letters opposing Secretary Casper Weinberger's "maximum allowable costs" plan aimed to cut Medicare and Medicaid drug costs. Opposition appears evenly divided the spokesman said, among pharmacists, M.D.s, and industrial and professional societies. M.D.s are chiefly concerned with questions of quality and interchangeability and interference with practice of medicine.

POSTPONED — Medicare-Medicaid utilization review requirements for hospitals and nursing homes are now set back to July 1 because many rural areas were unable to establish procedures for former Feb. 1 deadline. VINYL CL — New occupational standards that call for maximum exposure of 5 ppm of vinyl chloride are in effect pending appeal by manufacturers, after Supreme Court decision not to stay April 1 effective date for the standards.

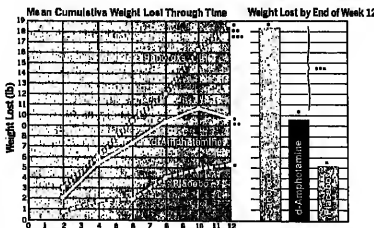


# SANOREX® IN OBESITY

(MAZINDOL)® TABLETS, 1 mg and 2 mg.

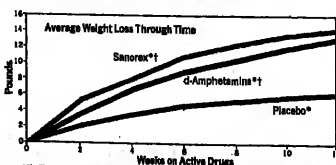
the soft underbelly  
of American health

## AS EFFECTIVE AS d-AMPHETAMINE

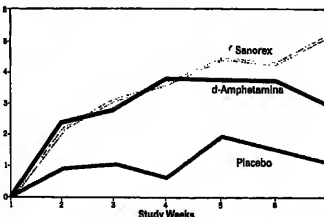


In a double-blind study of 40 obese patients (all of whom completed the study), Sanorex (1 mg t.i.d.) was more effective than either placebo or d-amphetamine (5 mg t.i.d.) in helping patients lose weight.

The 14 patients on Sanorex experienced a substantially greater mean weight loss—1½ to 2 lb/wk, as compared with 1 to 1½ lb/wk for the 14 d-amphetamine patients—throughout the 12-week phase of active medication. After the sixth week, the superiority of Sanorex became increasingly evident. And as treatment progressed, so did weight loss in patients on Sanorex—whereas after the tenth week, patients on d-amphetamine began to regain some weight.



\*Statistically significant weight loss from modified baseline (p < 0.05).  
†Statistically significant weight loss from placebo at each two-week interval (p < 0.05).



In a double-blind study of 90 obese patients (59 of whom completed the study), Sanorex (1 mg t.i.d.) was more effective than either placebo or d-amphetamine (5 mg t.i.d.) in helping patients lose weight.

By the end of the third week of active medication, weight loss in the 20 d-amphetamine patients began to plateau, and by the end of the fifth week, these patients began to regain some weight. On the other hand, the 18 patients on Sanorex continued to lose weight throughout the six-week course of therapy.

In a double-blind study of 93 obese patients (all of whom completed the study), 30 patients received Sanorex (1 mg t.i.d.), 31 received placebo, and 32 received d-amphetamine (5 mg t.i.d.).

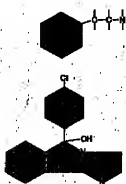
During the 12-week phase of active medication, patients on Sanorex lost an average of 14.1 lb, compared with 13.1 lb for d-amphetamine patients and 5.6 lb for placebo patients. Throughout the active medication phase, 63% of patients on Sanorex lost more than 1 lb/wk, compared with 38% of the d-amphetamine group and 29% of the placebo group.

## BUT WITH CERTAIN DIFFERENCES

Although the pharmacologic activity of Sanorex and that of amphetamines are similar in many ways (including central nervous system stimulation in humans and animals, as well as production

of stereotyped behavior in animals), animal experiments suggest that there are differences. Sanorex also differs in basic chemical structure from amphetamines and all other prescription anorexics.

### Different Chemical Structure



An important chemical similarity between amphetamines and all other prescription anorexics except Sanorex is the basic phenethylamine structure to which their differentiating chemical radicals are attached.

An important chemical difference between Sanorex and all other prescription anorexics is that Sanorex (as the isomer, it does not contain a phenethylamine structure.

### Different Neurochemical Action

**Action of d-Amphetamine** In animal studies, d-amphetamine (like intake of food) activates afferent neurons leading to appetite centers in the hypothalamus. Resulting release of norepinephrine activates the receptor neurons. Unlike food, however, d-amphetamine also suppresses norepinephrine synthesis. Thus, increasingly larger doses of d-amphetamine become necessary to produce an effect.\*

**Action of Sanorex (mazindol)** After intake of food stimulates the release of norepinephrine from the afferent neuron, Sanorex blocks its re-uptake without disturbing normal synthesis and release.\*

\*The significance of these differences for humans is uncertain.

### Simplicity and Flexibility of Dosage

Simple one-a-day dosage is facilitated by 2-mg tablets (taken 1 hour before lunch).

New flexibility (for the patient in whom 1 mg t.i.d. is preferred) is now facilitated by new 1-mg tablets (taken 1 hour before meals).

For Brief Summary, please see facing page.

Wednesday, April 16, 1975

## SANOREX® (MAZINDOL)®

**References**  
1. Kornhaber A. Problems and current concepts in the treatment of obesity. Scientific Exhibit presented at the New York State Academy of Family Physicians 22nd Annual Scientific Convention, October 10-11, 1973.  
2. Gaffney EA, Chaykin LB, Cohen AJ. Double-blind clinical evaluation of mazindol, dextroamphetamine, and placebo in treatment of morbid obesity. *Ann NY Acad Sci* 238: 107-119, 1975.  
3. Vercellotti JR. Practical considerations for managing obese patients. In: *Obesity: Scientific Exhibit presented at the American Medical Association, 27th Clinical Convention, Anaheim, Calif, Dec 1-4, 1973.*

Indication: In exogenous obesity, as a short-term (a few weeks) adjunct in a weight reduction regimen based on caloric restriction. The limited usefulness of agents of this class should be measured against possible risk factors.

**Contraindications:** Glaucoma; hypersensitivity or idiosyncrasy to the drug; established states; history of drug abuse; during, or within 14 days following, administration of monoamine oxidase inhibitors (hypertensive crisis may result).

**Drug Interactions:** May decrease the hypotensive effect of antihypertensive drugs; may develop within a few weeks; if this occurs, do not exceed recommended dose; do not combine with other sympathomimetic agents; such as operating machinery or driving a motor vehicle; and patient should be cautioned accordingly.

**Drug Dependence:** Mazindol shares important pharmacologic properties with amphetamines and related stimulant drugs that have been extensively abused and can produce tolerance and severe psychologic dependence. Manifestations of psychologic dependence or withdrawal with mazindol have not been determined in humans. Abstinence effects have been observed in dogs after abrupt cessation of prolonged periods. There was some self-administration of the drug in monkeys. EEG studies and clinical scores in human subjects yielded equivocal results. While the abuse potential of mazindol has not been further defined, possibility of dependence should be kept in mind when evaluating the desirability of including this drug in a weight-reduction program.

**Use in Pregnancy:** In rats and rabbits an increase in neonatal mortality and a possible increased incidence of foetal anomalies in rats were observed at relatively high doses. Although these studies have not indicated important adverse effects, the use of mazindol in pregnancy or in women who may become pregnant requires that potential benefit be weighed against possible hazard to mother and infant.

**Use in Children:** Not recommended for use in children under 12 years of age. **Precautions:** Insulin requirements in diabetic mellitus may be altered. Smallest amount of mazindol feasible should be prescribed or dispensed at one time to minimize possibility of overdose. Use cautiously in hypertension with monitoring of blood pressure; not recommended in severe hypertension or in symptomatic cardiovascular disease including arrhythmias.

**Adverse Reactions:** Most commonly, dryness of mouth, constipation, nervousness, and insomnia. Other reactions: Palpitation, tachycardia, Central Nervous System: Overstimulation, restlessness, dizziness, insomnia, tremor, hypotension, headache, depression, drowsiness, weakness, gastrointestinal distress of mouth, unpleasant taste, diarrhea, constipation, loss of appetite, other gastrointestinal disturbances.

**Side Effects:** Rash, excessive sweating, clamminess, endocrine imbalances. Changes in libido have rarely been observed. Eye: dogs resulted in severe corneal opacities reversible on cessation of medication; no such effect has been observed in humans. **Dosage and Administration:** 1 mg, three times daily, one hour before meals, or 2 mg per day, taken one hour before lunch as a single dose.

**Supplied:** Tablets, 1 mg and 2 mg, in packages of 100.

**Before prescribing or administering, see package insert for Prescribing Information.**

SANOREX PHARMACEUTICALS, EAST HANOVER, N.J. 07930

## Gene Engineering Work Gets Proceed-With-Caution Signal

By JUDITH RANDAL  
Special Tribune Correspondent

PACIFIC GROVE, CALIF.—Investigators who last summer voluntarily limited experiments in which bacteria—chiefly *Escherichia coli*—were fitted with foreign genes now have decided that the work can resume if good laboratory housekeeping practices, stringent personnel discipline, and other safety measures are rigorously observed.

The decision was reached during a four-day meeting here in which 86 American biologists and 53 from abroad came to grips with the possibility that DNA recombinants might accidentally unleash pathogenic unknown in nature and for which there would be no remedy.

### Outgoing Evolution

"The issue . . . [is that] a new technology of molecular biology appears to have allowed us to outdo the standard events of evolution by making combinations of genes which are unique in natural history," said Dr. David Baltimore of the Center for Cancer Research at the Massachusetts Institute of Technology in opening the conference.

While it was agreed that the experiments promise for the first time to offer practical and theoretical solutions to many problems in agriculture, biology, and medicine, there was also a consensus on the need for some strict controls.

The concern stems from the discovery about five years ago of "restriction" enzymes. These catalysts have enabled biologists to open up sequences of DNA from bacteria at known points and to insert into them equally well-defined DNA sequences from bacteriophages, mammalian or avian viruses, or eukaryotes such as fruit flies, mice, slime molds, sea urchins, and South African toads.

Because of the precision with which the enzymes cleave, the recipient DNA quickly lends after being grafted and is presumably capable of utilizing the new hereditary information both in the bacterium made to carry the hybrid molecules and their descendants.

### Infectivity Unpredictable

As far as is known, no harm has resulted from any of the experiments conducted to date. However, the infectivity of DNA recombinants is unpredictable and several investigators have expressed anxiety that pathogenicity might inadvertently spread to laboratory workers, the environment, and the public by any of several routes.

In addition to the DNA of their chromosomes, for example, bacteria often carry ringlets of DNA called plasmids which are readily exchangeable not only among *E. coli*, but also among *E. coli* and other species such as *Shigella* and *Streptococcus*. Since the plasmids are convenient vehicles for introducing foreign genes into bacteria and genes for antibiotics resistance have been used as "markers" in some recombinant experiments, the fear is that some novel form of drug resistance might be unleashed.

Similarly, combinations of bacterial

DNA and either viral or animal nucleic acids can be made so that the resulting hybrid molecules contain sequences in common with those of certain oncogenic viruses. About a year ago, for instance, the DNA of a human virus was linked to an RNA tumor virus of humans. Although the experiment stopped short of inserting the material into bacteria, the enzyme technique could have been used to make either plasmids or bacteriophages serve this purpose.

Thus, while such methods present investigators with unprecedented opportunities to understand the neoplastic process at the molecular level, they also pose the specter of possible carcinogen spread. And while cloning of DNA recombinant bacteria may well prove a new and abundant source of valuable medicinal materials, there is also the possibility that amplification of some concomitant and undesirable gene product might get out of hand.

It was against this background that the National Academy of Sciences and its operating arm, the National Research Council, appointed a special search committee of biologists last year. Chaired by Professor Paul Berg, a Stanford University biochemist, the committee rocked the scientific community in July when—in a letter published by *Science* and *Nature* magazines—it called for an international conference to consider the problems and meanwhile urged a voluntary world-wide moratorium on the potentially riskiest experiments.

### Unofficial Compliance

Although the ban was made official only in Britain, biologists from all the other nations represented at the meeting (Australia, Belgium, Canada, Denmark, France, Germany, Italy, the Netherlands, Japan, Poland, the Soviet Union, Switzerland, and the United States) appear to have complied. Speaking for the Soviet delegation, for example, Academician A. A. Bayev of the Institute of Molecular Biology in Moscow indicated that his country was gearing up to do recombinant research, but was delaying actually doing so until after the international conference met.

Similarly, a working party headed by Lord Ashby, Master of Clare College, Cambridge, found in its report to the British Medical Research Council in December that "the techniques open up exciting prospects both for science and education to society" and that "the potential hazards can be kept under control." But the M.R.C., like the U.S.S.R., planned to see what the international conference would do before lifting its ban.

In confronting the biohazards problem, the conference drew up a draft proposal whose principles will probably be followed by research funding agencies throughout the world.

Under the provisional guidelines, pending experiments would be classified by potential risk so that some could be performed under conditions that prevail in the typical university or hos-

pital microbiology laboratory, while others would require high security containment similar to that the astronauts experienced in quarantine after their return from the moon. And still others would be deferred indefinitely until new precautionary measures could be developed, tested and put in place.

The ultimate safety goal, however, is to alter the biological properties of the experimental materials themselves in such a way as to make it exceedingly unlikely that the test organisms could replicate should they escape from the laboratory and somehow find their way into humans.

### 'Fail-Safe' Strains Envisioned

Envisioned are "fail-safe" strains of bacteria—some already in existence—that cannot reproduce their kind except when supplied with certain crucial factors such as ultraviolet light, extremes of hot or cold, or special nutrients.

The idea would be to fit each strain of, say, *E. coli*, with a number of such inbred dependencies. According to Dr. Sydney Brenner of Britain's Medical Research Council, organisms could quickly be selectively created whose chances of replicating in nature would be as low as 10<sup>-24</sup> (one in a trillion). Non-transmissible plasmids and bacteriophages are among the other precautionary possibilities, as are alternatives to antibiotic resistance, such as "marker" genes.

The conference here was sponsored by the National Academy of Sciences and paid for by the National Institutes of Health and the National Science Foundation. The day after it closed in San Francisco to consider how the draft proposal might be translated into specific criteria for the awarding of contracts and grants in the United States. Among the possibilities considered was that researchers agree in advance to immediately shun any fail-safe biological materials they developed with the scientific community at large or free loss of funding support.



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## Hot Price Label Fumes Implicated in Meatwrapper's Asthma

Medical Tribune Report

SAN DIEGO, CALIF.—Fumes from hot meat price labels and not the polyvinyl chloride (PVC) in plastic wrappings have been implicated as the primary culprit in "meatwrapper's asthma," an occupational disease first reported about two years ago.

Within 30 seconds to twenty minutes after exposure to fumes produced when meat package labels were heated on a commercial labelling machine, nine of 13 meat-wrapping personnel developed immediate severe bronchoconstriction in a study, carried out under simulated working conditions, by Dr. Rudi Andrasch and his colleagues at the University of Oregon Health Science Center.

Eight workers developed paroxysmal cough and deep cyanosis, tachycardia, diaphoresis, and dizziness were observed in five others, Dr. Andrasch told the 31st annual meeting of the American Academy of Allergy here. In addition, other workers in the study reported skininess, severe burning in the nose and throat, headache, nausea, muscle ache, rhinitis, weakness, vomiting and hoarseness, he said.

### Response to Medication

"Three patients showed an excellent response to subcutaneous epinephrine, four patients required intravenous aminophylline in addition, and two patients continued to manifest severe bronchoconstriction requiring additional treatment with IPPB and phenylephrine with isocathrine.

"This study indicates that the fumes of thermocoated price labels are the principal culprit of meat-wrapper asthma. Paroxysmal cough and acute bronchoconstriction developed more frequently after shorter periods of exposure [than to PVC fumes] and tended to be much more severe.

"Dryness and burning of the mucous membranes, severe headache, extreme irritability and nausea were frequent

associated symptoms. Fumes of polyvinyl chloride soft wrap resins are also mucous membrane and respiratory irritants which may in a smaller group of meatwrappers cause moderate to severe reactive airway disease," Dr. Andrasch explained.

Although the fumes from the heated meat labels have still not been chemically identified, the adhesive backings are known to contain miscellaneous elastomers, thermoplastic copolymers, styrene butadiene copolymers, styrene acrylonitrile copolymers, polyphenylene oxides, polysulfones and phthalic acid plasticizers, the Oregon researcher said. Fumes produced when PVC wrappings are sliced by hot wire cutters, include carbon monoxide, carbon diox-

ide, hydrochloric acid, and various plasticizers, as well as hydrocarbons and chlorinated emulsions.

### Syndrome a Complex Response

"Even though price label fume intolerance accounts for most of the respiratory symptoms, the entire spectrum of the meat wrapper's syndrome has to be interpreted as a complex response to both PVC and price label fume exposure," Dr. Andrasch emphasized.

The Oregon study, supported by the local Meat Cutter's Union, followed a survey of 67 meatwrappers in the Portland area. Of those who responded, 57 per cent reported "moderate to severe respiratory symptoms," while a smaller number complained of head-

ache, sore throat, stomach cramps, and nasal congestion among other symptoms. In some cases, the attacks began three to four hours after the employee commenced work, but others began to have difficulties within the first ten to fifteen minutes on the job.

At the present time, two court cases involving meat wrapper's asthma are pending, one in Portland and one in Kentucky, Dr. Andrasch said, noting that from 25,000 to 50,000 meatwrappers, meat cutters, and supervisory personnel are exposed to fumes from PVC or label adhesives in their work.

New film cutting machines seem to be reducing the PVC hazard at this time, Dr. Andrasch concluded, suggesting that the hazards from meat label fumes could also be minimized through the expanded use of recently developed automatic labelling machines.

## after taking a potent analgesic 30 minutes ing



### how big a dose will now bring relief if it is a narcotic?

"Tolerance is an ever-present hazard to continued use of narcotics. . . . The very first dose diminishes the effects of subsequent doses." And, as increasing amounts of narcotics are required to control pain, distressing adverse effects—lathargy, hypotension, constipation, etc.—can needlessly debilitate the patient.

1. Sedation, M. S. A. Look at narcotic and non-narcotic analgesics, *Hospital Med.* 40:132, June 1971.

### how big a dose will now bring relief if it is Talwin?

Chances are, the same 50 mg. Talwin Tablet you prescribe originally will continue to provide good pain relief. Talwin can be compared to codeine in analgesic efficacy: one 50 mg. tablet appears equivalent in analgesic effect to 60 mg. (1 gr.) of codeine. However, patients receiving Talwin Tablets for prolonged periods face fewer of the consequences you've come to expect with narcotics. There should be fewer "adverse effects" on her way of life.

Tolerance rare: Tolerance to the analgesic effect of Talwin Tablets is rare.

Dependence rare: During three years of wide clinical use, there have been a few reports of dependence and of withdrawal symptoms with orally administered Talwin. Patients with a history of drug dependence should be under close supervision while receiving Talwin orally.

In prescribing Talwin for chronic use, the physician should take precautions to avoid increases in dose by the patient and to prevent the use of the drug in anticipation of pain rather than for the relief of pain.

Generally well tolerated by most patients: Infrequently causes decrease in blood pressure or tachycardia, rarely causes respiratory depression or urinary retention, seldom causes dizziness or constipation. Acute, Autonomic CNS effects, described in product information, have occurred in rare instances following the use of Talwin Tablets. If dizziness, lightheadedness, nausea, or vomiting is encountered, these effects may decrease or disappear after the first few doses.

\*See important product information for adverse reactions, patient selection, prescribing and precautionary recommendations.

### in chronic pain of moderate to severe intensity

**Talwin<sup>®</sup> 50 mg. Tablets**  
brand of  
**pentazocine**  
(as hydrochloride)

Talwin is not subject to narcotic controls.

New suggested: Tablets, white, oval, scored. Each tablet contains Talwin (pentazocine) as hydrochloride equivalent to 50 mg. base, Codeine 100.

Wednesday, April 16, 1975

## Ship-to-Shore Electrocardiography Initiated



Zim company, the Israeli shipping line, has installed ship-to-shore electrocardiography on one of its vessels in what is believed to be the first trial of its kind. The machine is connected by radiotelephone to a monitor at Rambam Hospital in Haifa. The initial test, carried out when the ship was 600 miles off Haifa, was reported to have produced surprisingly clear results.

MEDICAL TRIBUNE

## California Acupuncture Body With Few MDs Expected Soon

By EDWARD GROSSMAN  
Medical Tribune Staff

SACRAMENTO, CALIF.—Despite opposition from some physicians, it seems likely that California will soon have a law creating an Acupuncture Advisory Board, comprised of two M.D.s and five "traditional" acupuncturists, which would be empowered to issue certificates of qualification to practice acupuncture as a healing art.

Bill number 86, introduced by Senator George Moscone, D-San Francisco, was recently passed by a vote of 22-1 in the State Senate. John Jervis, aide to Sen. Moscone, told MEDICAL TRIBUNE that he expects smooth sail-

ing in the Assembly as well. The bill would then have to be signed by Gov. Edmund Brown, Jr., and would take effect as law Jan. 1, 1976.

A similar bill was approved overwhelmingly by both houses of the legislature last September, but was vetoed by former Gov. Reagan, who said that acupuncture was still in the research stage.

### Restriction Floated

Under present California law, only licensed professionals (M.D.s, D.D.S.s, and chiropractors) are permitted to puncture the skin for therapeutic purposes. However, many non-licensed herbalists and "doctors of Chinese Medicine" have been practicing openly, with only sporadic prosecution and, infrequently, the imposition of small fines.

The Moscone measure would make it a misdemeanor to practice acupuncture or "hold oneself out as an acupuncturist" without a certificate issued by the new Advisory Board. Furthermore, acupuncture could not be performed without "prior diagnosis or referral" from a physician.

The Advisory Board is to be under the jurisdiction of the state Board of Medical Examiners and its members are to be appointed by the Governor. It will be charged with establishing standards, tests, and other requirements, and will pass on the applications of all licensure candidates, with acupuncturists having more than five years experience receiving priority consideration.

The law would not affect the right of physicians and dentists to practice acupuncture.

Several California physicians who find fault with the Moscone proposal concede that it is virtually assured of passage. Their main objection concerns the composition of the Advisory Board.

### 'Many Poorly Trained'

"There are many so-called 'traditional' acupuncturists, trained in Hong Kong and elsewhere, who are skilled and conscientious," Dr. Jane F. Lee, a San Francisco general practitioner, told MEDICAL TRIBUNE. "But there are also many others who have been poorly trained, and have little knowledge of physiology, pathology, anatomy. I'm worried that with the drastic imbalance on the Board, there might be wholesale certification of unqualified practitioners. I'd like to see a more balanced approach."

Dr. Lee is a member of the Acupuncture Committee of the California Medical Association, which so far has not taken a position on the bill.

Her view is shared by Dr. George Wong, Jr., a family practitioner on Long Beach, who, like Dr. Lee, has extensive training in acupuncture and uses it as an "adjunct mode" in his practice. Dr. Wong is Chairman of the Acupuncture Research Institute Alumni Association, a non-profit organization of physicians interested in ac-

Continued on page 23



## What a difference a day can make

Your counsel and reassurance—and Ritalin.  
A logical first step in treating mild depression and often all that's needed to bring quick symptomatic relief.  
Indeed, your patient may be-

gin to feel better within hours—her spirits boosted, her mood brightened. A single prescription may be all that's needed.  
Ritalin is usually well tolerated even by older or convalescent patients. Note, however,

that it is not indicated in the more severe depressions. But whenever depression is mild, think of Ritalin—so your patient has a better chance of waking up to a brighter tomorrow.

**Ritalin**  
(methylphenidate)  
acts quickly to relieve symptoms  
in mild depression

\*This drug has been evaluated as possibly effective for this indication. See label for complete information.

**Ritalin® hydrochloride (C)**  
(methylphenidate hydrochloride)  
TABLETS

**INDICATION**  
Based on a review of this drug by the National Academy of Sciences-National Research Council and/or other information, FDA has classified the indication as follows:  
"Possibly" effective: Mild depression  
Final classification of the less-than-effective indication requires further investigation.

**CONTRAINDICATIONS**  
Marked anxiety, tension, and agitation, since Ritalin may aggravate these symptoms. Also contraindicated in patients known to be hypersensitive to the drug and in patients with glaucoma.

**WARNINGS**  
Ritalin should not be used in children under six years, since safety and efficacy in this age group have not been established.  
Sufficient data on safety and efficacy of long-term use of Ritalin in children with minimal brain dysfunction are not yet available. Although a causal relationship has not been established, suppression of growth (ie, weight gain and/or height) has been reported with long-term use of stimulants in children. Therefore, children requiring long-term therapy should be carefully monitored.

Ritalin should not be used for severe depression or either exogenous or endogenous origin or for the prevention of normal fatigue states.  
Ritalin may lower the convulsive threshold in patients with or without prior seizures with or without prior EEG abnormalities, even in absence of seizures. Safe concurrent use with anticonvulsants and Ritalin has not been established. If seizures occur, Ritalin should be discontinued. Use cautiously in patients with hypertension. Blood pressure should be monitored at appropriate intervals in all patients taking Ritalin, especially those with hypertension.

**Drug Interactions**  
Ritalin may decrease the hypotensive effect of guanethidine. Use cautiously with other CNS agents and MAO inhibitors. Ritalin may inhibit the metabolism of coumarin anticoagulants, anticholinergics (parasympathetic blockade), tricyclic antidepressants (imipramine, desipramine). Downward dosage adjustment of these drugs may be required when given concomitantly with Ritalin.

**Usage in Pregnancy**  
Adverse animal reproduction studies to establish use of Ritalin during pregnancy have not been conducted. Therefore, no further information is available. Ritalin should not be prescribed for women of childbearing age unless, in the opinion of the physician, the potential benefits outweigh the possible risks.

**Drug Dependence**  
Ritalin should be given cautiously to emotionally unstable patients, such as those with a history of drug dependence or alcoholism, because such patients may increase dosage on their own initiative.

Chronic abuse of Ritalin can lead to marked tolerance and psychic dependence with varying degrees of abnormal behavior. Frank psychic dependence can occur, especially with parenteral abuse. Careful supervision is required during drug withdrawal, since severe depression as well as the effects of chronic overactivity can be manifested. Long-term therapy should be required because of the patient's basic personality disturbances.

**PRECAUTIONS**  
Patients with an element of agitation may react adversely to discontinuation of therapy. If necessary, barbiturate, chloralhydrate, and phenobarbital are advised during prolonged therapy.

**ADVERSE REACTIONS**  
Nervousness and insomnia are the most common adverse reactions but are usually controlled by reducing dosage and continuing the drug in the afternoon or evening. Other reactions include: hyperemesis, tachycardia, dry mouth, urinary retention, arthralgia, exfoliative dermatitis, arthralgia, multiple myeloma, histiocytosis, findings of necrotizing vasculitis, and thrombocytopenic purpura; anorexia, weight loss, dizziness, vertigo, headache, dyslexia, drowsiness, blood pressure and pulse changes, both up and down, tachycardia; angina, cardiac arrhythmias; abdominal pain; weight loss during prolonged therapy. Toxic psychosis has been reported. Although a causal relationship has not been established, the following have been reported in patients taking this drug: leukopenia and/or anemia; a few instances of scalp hair loss.

In children, loss of appetite, abdominal pain, weight loss during prolonged therapy. In adults, tachycardia may occur more frequently. However, any of the above adverse reactions listed above may also occur.

**DOSEAGE AND ADMINISTRATION**  
Adults  
Administer orally in divided doses 2 or 3 times daily. The usual dosage is 20 to 30 mg daily. The maximum daily dose is 60 mg. The minimum daily dose is 10 mg. The minimum daily dose is 10 mg. The minimum daily dose is 10 mg.

**HOW SUPPLIED**  
Tablets, 20 mg (pink, scored), bottles of 100 and 500.  
Tablets, 10 mg (pink, scored), bottles of 100, 500, 1000 and 2000. Also 10 mg (pink, scored), bottles of 100, 500 and 1000.

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**C I B A**

Wednesday, April 16, 1975

MEDICAL TRIBUNE

11

The Only Independent Weekly Medical Newspaper in the U.S.

# Medical Tribune

and Medical News  
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## Genetic Engineering

THE INTERNATIONAL CONFERENCE ON Recombinant DNA Molecules, which met in late February in Asilomar, Calif., has drawn up a list of recommendations on precautions to be taken by investigators working in the field of genetic engineering (see page 5). Technical skills now available make it possible with the use of certain enzymes to cleave DNA at specific sites and join DNA from animal viruses with bacterial DNA or with viral DNA. The possibilities include illumination of the very basics of gene action.

But a year and a half ago, the potential hazards of "new kinds of hybrid plasmids or viruses, with biological activity of unpredictable nature" was raised at the 1973 Gordon Conference on Nucleic Acids. A Committee on Recombinant DNA Molecules, chaired by Professor Paul Berg of the biochemistry department at Stanford University, was formed in 1974 and in July called for a voluntary suspension of certain types

of genetic manipulation until a conference of workers in the field could be held to spell out precautions and tabus.

The precautions discussed at the conference include the high skills of the investigators themselves, the careful laboratory practices needed, and the use of biological barriers, such as organisms capable of survival only in the special environments of laboratories and not in natural environments.

Nobel Laureate Joshua Lederberg has expressed the fear that safeguards and precautions "that are entirely appropriate for certain risks might be prematurely rigidified into a set of bureaucratic regulations that might be very readily enforced beyond the dominion of their reasonable application." That is a fair warning, the precautions and the safeguards are in the hands of the investigators themselves. It is up to them to keep control in their own hands and out of the hands of bureaucrats.

## Solving the Riddle of Diabetes Mellitus

IT IS NOW 54 years since Banting and Best demonstrated that an extract of the islet tissue of the pancreas can lower the blood sugar of the diabetic dog. Although the successful preparation of insulin extracts seemed at first to have solved the riddle of diabetes mellitus, this enthusiastic belief soon waned. Indeed, in the past quarter of a century, the puzzling questions about the etiology and pathogenesis of diabetes have multiplied rather than diminished as the techniques for investigating the disease have become more sophisticated and precise.

There are many reasons why the definition of diabetes mellitus as simply a disorder resulting from a relative or absolute deficiency of insulin secreted by the beta cells of the pancreas is unsatisfactory. For the past several years, Dr. Roger H. Unger of the Veterans Administration Hospital in Dallas, Texas, and the U. of Texas Southwestern Medical School, has championed the notion that the disease is a bimodal disease, "which holds that the major consequence of absolute or relative insulin lack is glucose underutilization and that absolute or relative glucagon excess is the principle factor in the overproduction of glucose in diabetes."

There is powerful evidence in support of this concept. The alpha cells of the pancreas secrete glucagon, which is

known to be a hyperglycemic hormone. There has been ample demonstration since the late 1960's that every form of diabetic and non-diabetic hyperglycemia investigated "is accompanied by relative or absolute hyperglucagonemia."

Since the discovery of somatostatin, the hypothalamic growth hormone-release-inhibiting factor, it has been found to suppress both glucagon and insulin secretion. A series of brilliant investigations in Dr. Unger's laboratory and in a number of independent laboratories have demonstrated that hyperglycemia in dogs made insulin-deficient by alloxan or total pancreatectomy is abolished by somatostatin injection.

Dr. John E. Gerich and his colleagues at Dr. Peter Forsham's laboratory at the U. of California in San Francisco have recently reported that somatostatin injection in insulin-dependent diabetes reduced their plasma glucagon and hyperglycemia as well. As the investigators state: "The present findings have important therapeutic implications." What is needed is a preparation of somatostatin with a prolonged half-life. The exciting and intriguing possibility is that failure to prevent microangiopathy and atherosclerosis in diabetes by insulin therapy may be turned to success with the addition of somatostatin to the therapeutic regimen. Time will tell.

## Hepatitis B Virus Carriers

CLINICAL QUOTE: "The significance of hepatitis B infection in early life lies also in its importance in the presence of prolonged carriage of hepatitis B virus. Zuckerman and Taylor (1969) described a well-documented healthy former blood donor carrying

hepatitis B antigen for at least 20 years. That a reservoir of chronic carriers may become established among children is, therefore, a cause for the utmost concern." (Dr. Arie Zuckerman, at a March of Dimes-National Foundation symposium on infections.



"I'll tell you something else I'm learning to live with!  
Doctors who say, 'Learn to live with it!'"  
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## LETTERS TO TRIBUNE

### Carotid Artery Palpation

I must take exception to one statement of Dr. Edwin Beven's (MT, Mar. 5).

Noninvasive or not—any palpation, compression, movement of adjacent structures, etc.—concerning a carotid artery with even minor stenosis cannot be considered a "No-Risk Method."

While the risk may be slight, the possibility of causing sudden, complete occlusion or cerebral embolization does exist and should be taken into consideration before even getting close to a patient's carotid artery. The presence of anomalous circulation, stenosis of other vessels, elevated lipoproteins, prior TIAs and a variety of other factors will, of course, increase the risk.

DONALD M. POSNER, M.D.  
CNSAR, VT.

administered, that patient will damned well be his own watchdog to see that his doctor's orders are carried out to the letter.

I personally have, in two highly respected medical institutions, refused attempted treatments which I knew were not ordered by my physician. Since I have the benefit of having been a medical journalist for 15 years I knew one procedure would have resulted in a fast trip to the intensive care unit, if not the morgue.

As doctors you owe that to your patients. Tell them to yell their heads off if anyone attempts treatment counter to your instructions. Who knows? If that were made standard procedure there might be a marked drop in malpractice suits.

HARRY WELKEN  
New York

### New Laws Needed?

Dr. David Nathan is quoted (MT, Feb. 12) as follows: "I never occurred to us that society would be worried that we will not maintain life." This must certainly strike a new low in what used to be called a "life science" of medicine. Poor Dr. Nathan is so preoccupied with what he sees on the other end of his microscope that he forgets that he is in the business of saving lives.

Irresponsible ivory tower pseudoscientists who have used unborn children as lab animals in vivisection-type experiments have created the need for new laws. They would like to use "hostility to abortion" as their whipping boy but the real culprit is their own ignorance of the Declaration of Helsinki.

THOMAS J. BOAN, M.D.  
Chicago

### Endorsing Euthanasia?

The American Association of Pro Life Obstetricians and Gynecologists, comprised of members of our specialty from throughout the entire United States, wishes to make some observations regarding the Edelin verdict.

The Supreme Court decision of January, 1973 removed abortion from the criminal code and put it outside the law by making the mother and her physician entirely responsible for the destruction of human life. No mention was made in the Supreme Court decision as to what happens to the infant born alive and struggling for survival. As there have been hundreds, and possibly thousands, of these sadly unfortunate infant victims, sound medical practice and compassion should motivate all of us to render these babies the best possible medical care for survival.

As obstetricians and gynecologists, we have all seen infants of less than two pounds birth weight survive and do well when given the proper care. Why shouldn't we feel obligated to render the same care to the survivor of a miscalculated abortion procedure that we render to the infant in a normal delivery?

If the obstetrician has the right to destroy the live-born infant to an abortion procedure, would he not have the same right to extinguish the life of a newborn infant with a congenital defect whose mother may not want him?

The acceptance of this principle surely would, in fact, be providing a legal endorsement of euthanasia.

MATTHEW J. BULFIN, M.D.  
President  
Lauderdale-By-The-Sea, Fla.

### Telling the Patient

Right on for Dr. Eli Friedman "Panels Disagree on How Much To Tell Patient" (MT, Mar. 12).

For the moment let us lay aside the complex and tricky question of whether or not to inform a patient if he or she is dying. Rather, let us consider the non-comatose patient of average intelligence. Even in the best run hospitals mistakes are made, orders scribbled illegibly on a patient's chart, residents and nurses carelessly or too hastily briefed on medications and/or general management of a case.

If the physician will take the time to explain carefully to the patient what he is being treated for, what the medication is, and how and when it should be



## Nitroglycerin Reported 'Consistently Beneficial' in Infarction

Continued from page 1

sign of hypotension and reflex tachycardia they have also administered the vasoconstrictor phenylephrine to mitigate these two side effects.

In general, Dr. Epstein noted, M.I. patients fall into two subgroups. One consists of those in heart failure, with elevated pressures and inadequate pumping action. "In this group, nitroglycerin appears to be very effective in reversing the manifestations of heart failure," Dr. Epstein said. "The high pressures that build up in the lungs decrease to normal very abruptly after nitroglycerin administration and the heart starts pumping more effectively. That's not new; these effects have been demonstrated by groups at Cedars of

Lebanon Hospital in Los Angeles, at Massachusetts General Hospital, and at Johns Hopkins. But what we have found in addition is that the size of the infarct, the amount of muscle damaged, is significantly reduced by treatment with nitroglycerin."

### Non-Heart-Failure Group

The second subgroup of patients are those who have suffered a heart attack, Dr. Epstein continued, who have damaged muscle, but are not in heart failure. These patients are not usually treated, he said, "but nevertheless about 10 per cent of them die in hospital and another 10 per cent die within a year, so it is not a benign disease. In this subgroup of patients who have never been

treated before, we found exactly the same thing. We didn't get them out of failure, because they weren't in failure to begin with, but we found that the combination of nitroglycerin and phenylephrine reduced infarct size."

Dr. Epstein noted also that it is the second subgroup of patients that most often requires the phenylephrine and nitroglycerin. And in the future, he said, he and his associates plan to give the nitroglycerin intravenously so it can be monitored more readily and more precisely.

On the basis of their experience with animals, Dr. Epstein said he thinks the nitroglycerin exerts its beneficial effects in two ways: by increasing the amount of blood delivered to the ischemic area

via the collateral system, and by reducing the size of the heart chamber, thus decreasing myocardial tension and oxygen requirements.

Dr. Epstein is working with Dr. Kenneth M. Kent, Dr. Robert E. Goldstein, and Dr. David R. Redwood, of the N.H.L.L., Cardiology Branch, and Drs. Harrie Levitt and Norman Cagin, of New York, Dr. Jeffrey S. Borer, who is in London on sabbatical leave from the N.H.L.L., is also participating in the trials.

Whether the treatment increases long-term survival is yet to be ascertained, Dr. Epstein said. But he thinks it has "enormous potential" and that long-term studies should be mounted. Dr. Epstein's animal studies are described in the January 2, 1975, *New England Journal of Medicine*, and preliminary clinical data appear in the April Journal of *Clinical Investigation*. Dr. Borer will present the full clinical report to the American Society of Clinical Investigation in May.

## Hepatitis B Virus Pool in Newborn Seen Building Up

Continued from page 1

and sponsored by the National Foundation-March of Dimes.

An evaluation of recent studies makes it clear, he emphasized, that both transplacental and perinatal transmission of hepatitis B infection from mother to child may take place in spite of older notions to the contrary.

Dr. Zuckerman recommends that all antigen-positive mothers be instructed to pay scrupulous attention to personal hygiene when handling their infants. Since there is the possibility of transmission of the antigen via breast milk, he believes breast feeding in these cases should be discouraged.

### Control Attempted in Newborn

Control of hepatitis B infection in the newborn is now being attempted by passive immunization with specific hepatitis B immunoglobulin and by immunotherapy with transfer factor, he commented, adding that safe and effective hepatitis vaccines—"now under development"—are a pressing need.

The limited data available indicate that the frequency of transmission of hepatitis B from mother to infant is highest (76 per cent in one study) when acute infection occurs during the third trimester of pregnancy or early in the postpartum period, and relatively low (10 per cent) if it develops during the first six months, Dr. Zuckerman said.

Investigations of transmission by asymptomatic carrier mothers have yielded variable figures but in one Japanese study cited by Dr. Zuckerman eight of 11 infants born to such mothers showed antigen in their sera within six months of delivery and the antigen persisted during prolonged follow-up.

Many of the infants in whom antigen is detected remain clinically well, the virologist said, although some show "prolonged elevation of an enzyme frequently associated with liver damage."

## Control Program Credited With 10% Decrease in Hospital Infections

Continued from page 1

The study, which was conducted at the Presbyterian-St. Luke's Hospital between 1969 and 1973 and computer analyzed 11,656 occurrences of nosocomial infection, revealed a 10 per cent decrease in hospital infections at a time when more patients susceptible to infection were being admitted, said Dr. Edwards, who was the epidemiologist at the 840-bed hospital during the study period.

"Previous thorough studies have been very short-term and thus hard to interpret because there may be fluctuation in hospital infection rates from month to month," he said.

### Nearly \$850,000 a Year Saving

Attributing the decrease to the hospital's active infection control program begun in 1968, Dr. Edwards stated that the program generated a significant reduction in economic morbidity amounting to nearly \$850,000 per year, in addition to saving lives.

"We calculated this figure in the following way," he said. "The average patient stay in the hospital was 11-12 days during the study, varying from year to year. The average stay for patients with hospital onset infections, on the other hand, was 33 days, or an additional 21 days, of an average per diem charge of \$150. Considering that we encountered an average of 2,009 patients a year with hospital onset infections and multiplying these figures out, we estimate that the cost to patients and third-party payers for such infections was around \$6,328,350 a year. A 10 per cent reduction in infection rate, therefore, means an average savings of \$630,830 per year. Since the total cost of running our infection control program—including paying the salaries of three full-time nurse-epidemiologists and one half-time physician, as well as the costs of computerization—was around \$75,000 a year, the economic advantage of the program was considerable."

### Reduction in Mortality

The study also revealed significant findings with respect to mortality. "One can place patients dying of infection into three categories. In the first, the infection is the primary cause of death; in the second, the underlying disease is primary; and in the third, the infection contributes to but is not entirely responsible for the death. In our study we found that infection was the primary cause in 12.9 per cent of cases, that infection was an associated and contributing cause in 25.9 per cent, and that the underlying disease was primary in 61.2 per cent."

"Thus we concur with the literature that the underlying disease is determining in most cases, but at the same we concluded that if one can prevent some of the infections in the 12.9 per cent category, then one can reduce mortality as well as morbidity."

Looking at hospital onset infections in the context of total hospital mortality, Dr. Edwards indicated that about a fourth of patients who died at Presbyterian-St. Luke's between 1969 and 1973 had an infection of hospital origin at the time of death.

The epidemiological methods that eventually brought about the 10 per cent reduction in hospital onset infections at St. Luke's were scrutinized both during the formal study period and during a brief pre-computer pilot study in 1969. "In the earlier study we wanted to validate the effectiveness of the nurse-epidemiologists since many physicians at the time doubted their ability to accurately collect data. In comparing three fellows in the infections disease section with two nurse-epidemiologists over a two week period we found that the latter were 94 per cent as effective as the former in collecting and classifying data on infections, which is not a statistically significant difference. We specifically restudied this question several times throughout the next four years and found that the nurses were consistently as accurate as the physicians trained in infectious diseases."

"In the earlier study," Dr. Edwards added, "we also wanted to find out how many nurses are needed per number of beds to do the job and how often they need to visit the wards. Basically we concluded that one nurse was required for 300 beds and that they needed to visit the wards twice a week. After it was decided that we needed three nurses for our 840 beds, we hired an additional nurse."

### Combined Approach Used

Once a full-blown infection control program was launched at Presbyterian-St. Luke's in 1969, a combined epidemiological and teaching approach was followed. "There are four different approaches that hospitals may take," commented Dr. Edwards. "First, many hospitals merely have a perfunctory infection control committee that meets in order to fulfill accreditation requirements but doesn't really do anything about the endemic level of hospital infections and only becomes active if there is an outbreak. Unfortunately I suspect that this is the most common approach. Second, there is the command approach, in favor of which physicians will argue that we already know what most of the problems are and so let's go out and work on these and also be ready to investigate any epidemics that may come up. Unquestionably that approach will lower the infection rate at some sites at some hospitals, depending on the interests of the people who are commanding the commands, as it were. But it doesn't tell much about whether one is getting a total impact and a uniformly educated hospital staff to reduce the overall problem over the long haul."

"The third approach is what one might call the surveillance approach and implies simply collecting data. This has been much maligned because people should not just collect data and do nothing about it. The fourth approach, and the one we opted for, is the epidemiological one in which you do essentially all of the things that the other approaches do plus actively inform and instruct."

"Something I've always felt strongly about with respect to hospital infections," Dr. Edwards continued, "is that since we have such a large turnover

## Rate of Community Onset Infections and Hospital Onset Infections in Deaths By Service From 1969 Through 1972

Service	Rate of Infection in Death		Rate of Infection in Death		Underlying Disease Primary Cause	
	Primary Cause	HOI*	COI	HOI	COI	HOI
Medicine	14.3	8.4	25.2	20.0	6.2	40.2
Surgery	24.5	7.7	18.7	22.0	6.4	35.3
Pediatrics	24.5	2.5	80.9	15.3	2.7	55.8
Newborns	0.0	15.7	33.8	33.3	0.0	59.0
Total Hospital	15.3	12.9	24.8	25.0	6.0	37.2

HOI = Hospital Onset Infection  
COI = Community Onset Infection  
\*COI = Community Onset Infection (HOI = Hospital Onset Infection)

## Mortality Associated With Community Onset Infections and Hospital Onset Infections By Service From 1969 Through 1972

Service	Per Cent of Admissions		Per Cent of Deaths		Number of Infections Per Death	
	COI	HOI*	COI	HOI	COI	HOI
Medicine	1.7	1.7	22.3	22.3	1.4	1.2
Surgery	0.4	0.4	22.1	25.0	1.2	1.2
Pediatrics	0.0	0.0	1.9	8.3	0.0	0.0
Newborns	0.0	0.0	14.4	26.4	0.0	0.0
Total Hospital	0.8	0.8	14.4	26.4	0.0	0.0

COI = Community Onset Infection  
HOI = Hospital Onset Infection  
\*HOI = Hospital Onset Infection (COI = Community Onset Infection)

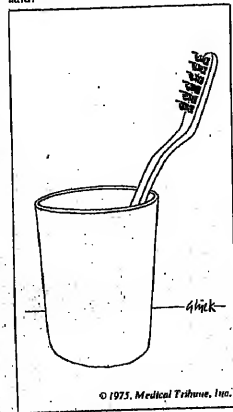
rate of hospital employees today, the key to control isn't so much new knowledge as it is having some ongoing way of continually bringing the problem to people's attention, both in terms of education about how to perform certain procedures and in terms of epidemiological data so one can know whether one is making an impact. If we're not, then push harder in the particular area where we're not making an impact. I don't think one can make reasonable applications unless one knows what is going on in one's own hospital."

### 80% of Infections at 4 Sites

In the Presbyterian-St. Luke's study it was determined that about 80 per cent of the infections were occurring at four major sites—the urinary tract, lower respiratory tract, surgical wounds and bloodstream. "All of the sites had pretty much the same decrease in infections except for the bloodstream, which actually went up. In 1969 there were 969 urinary tract occurrences compared to 741 in 1972, 665 lower respiratory tract occurrences compared to 651, 520 surgical wound infections compared to 402, and 173 bacteremias compared to 233. The rise in bacteremias, I think, may be due to the fact that the hospital started doing more bowel cancer surgery."

An unusual aspect of the study, and one which requires further investigation, is that it was the first long-term study to look at the interchange between community onset and hospital onset infections. "We defined a community onset infection as one present on admission or coming up to the first 72 hours and not related to a hospital procedure. I don't think we have any hard and fast data on this interchange, but we got some linking into where we need to look

further. For example, it's been well known that hospital personnel tend to have higher carriage rates of organisms like *Pneumococcus* compared to people in the community. Well, we counted several such classical infections that appeared to have their onset in the hospital; so the question is whether there is some interplay going on here that allows for spread of these organisms in the hospital at a greater frequency. We don't know much at all about how viral infections are introduced into and then spread throughout the hospital, and I think we can expect to see a greater research effort in this area," Dr. Edwards said.



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## Interdisciplinary PG Training Program Set

By MICHAEL HERRING  
Medical Tribune Staff

ROCHESTER, N.Y.—An interdisciplinary postgraduate training program at Strong Memorial Hospital utilizing internists, pediatricians, and specialists in the same physical setting—and in some cases operating as a fee-for-service private group practice—will go into full effect as soon as all services are moved into the new ambulatory care wing here, Dr. Warren Glaser, program coordinator, told MEDICAL TRIBUNE.

The program will provide for training at all levels, he said, and make physician services more readily available to all patients in the community, regardless of their ability to pay.

### Effective Coordination Sought

"We can't sacrifice the expertise we've gained from so much specialization, but at the same time, there has to be a way to coordinate individual efforts more effectively," he explained. "The beauty of the arrangement at Rochester is that you have both the generalist and the specialist working compatibly and in close proximity."

Dr. Glaser, who is Professor of Medicine and coordinator of ambulatory care, Department of Medicine at the University of Rochester School of Medicine and Dentistry, said that the decision to form the hospital team of group-practice internists and pediatricians, with the backup of subspecialists in each major area, was based on the recognition that "the hospital stands at the center of the ambulatory health care system."

"Ideally," he said, "each person in the community should have access to a personal physician who renders comprehensive care with continuity and who can delegate that care when necessary to the appropriate specialist."

Primary care, he continued, should be medical attention that is "available and accessible" to the patient when he or she needs it. "Primary care should not refer only to the initial visit to a doctor during office hours, but includes empathy, continuity, and treatment that is appropriate to the patient's changing needs," he said.

### Integrationist for Subspecialties

"At the same time, it should function to take the load off the emergency-room physician. Finally, the primary care physician is the integrationist for all medical subspecialties that the patient may require."

Dr. Glaser emphasized that the primary care physician at Strong Memorial will function increasingly as a member of a team—"not just with other doctors, but with nurses, social workers, and other medical and paramedical personnel."

"We think that the group practice of general internists and general pediatricians has more appeal," he commented, "because it is a higher level of care, and permits more appropriate referrals within the system."

Dr. Glaser briefly described the new arrangement at Strong Memorial as follows:

"In medicine, we have two or three interns, two assistant residents, and

two associate residents (team) with an attending physician and a licensed practical nurse. The whole team cares for a panel of patients."

In addition, he explained, the interdisciplinary group, together with the house staff, care for the medical clinic and combined clinic patients from the previous arrangement.

"These patients are now considered as one group of hospital patients, and are seen on a private-practice basis. Once a patient is entered into the system, the fees for hospital services and the fee-for-service practice are the same. Patients, no matter how they pay, can be transferred from one group to another. The only difference is in who does the billing."

The internal medicine group is a fee-for-service, private practice group, Dr. Glaser added. "These physicians cover for one another on a team basis."

"The doctor's offices will be in the hospital itself. If hospital patients require care after clinic hours, we use the emergency room. But we don't descend on the emergency room just because we can't provide care elsewhere."

### ER Resident on Call

"Rather, the resident in the emergency room acts as the on-call physician in a manner similar to those in the internal medicine group covering for one another. Obviously, we can't know all the patients, but we can provide care, based on the fact that this is a recog-

nized individual who has a problem. With all the groups working closely together, we have records and protocols on which to base a judgment."

"The continuity clinic is the patients' counterpart to the house staff group he added, but with a somewhat different organization. While the latter has interns, assistant residents, and associate residents working together, the pediatricians group is a horizontal arrangement, with all interns, all first-year residents, and all second-year residents working together, Dr. Glaser said.

He also pointed out that "patients here are actually participating in the practice, and medical students are free to view their work first-hand and form their own judgment. Naturally, we want it to be good so that the most able, interesting physicians of the future be attracted to this kind of patient care."

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MEDICAL TRIBUNE

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## One Man...and Medicine

ARTHUR M. SACKLER, M.D.  
Inventor of Dalmane, flurazepam HCl



### Now, A Word From the Opposition

MEDICAL TRIBUNE supports the free exchange of differing views. The following letters are some of the responses to Dr. Sackler's column on the Edelin case, "Doctor, Are You Innocent?" (MT, Mar. 12).—Ed.

Your newspaper's coverage of the Edelin conviction is worthy of the 1984 Blake is White Literary Award. The "straight" news story by a Special Cor-

respondent sets the tone in the headline—Shock and Dismay—with the predictable bias in the body of the story. Flanking this, we find a "Special Trib-

une Report" by the same author and your sterling piece, "Doctor—Are You Innocent?"

You and your newspaper seem somehow to have missed a basic point. A jury of his peers found Dr. Edelin guilty of manslaughter. He did not kill a fetus—he killed a living human being. Please remove my name at once from your mailing list.

WILLIAM DANIEL DAVIES, M.D.  
Evanston, Ill.

I was amazed and disappointed that a fine publication such as MEDICAL TRIBUNE dignified the likes of a Dr. Kenneth C. Edelin with a photo on your front page of the March 12, 1975 issue. Continuation of such published items only aid and abets the act committed.

Certainly there must be other topics

more interesting and deserving to sustain the name of good medicine which is being evaded daily by the very act about which he brags. Let us have no more of this, please. Thank you.

Incidentally, the editorial by Arthur M. Sackler, M.D., was about as enlightening as an overflowing commode.

GERARD A. DEL GRUPPO, M.D.  
Lock Haven, Pa.

The vile and vicious anti-Catholic tone of your editorial leads me to make this protest of your appeal to the worst instincts of the society. The kind of abortion performed by Dr. Edelin is disapproved by all segments of the society, all religions, and even a majority of atheists (see Blake, J. Science, April 1972).

The tortured non-sensitization of your argumentation lead me to believe that you were blinded by bigotry in departing from your usual well-reasoned rationale. You find it incomprehensible that a man could be found guilty of manslaughter in "standing by and denying a fetus oxygen and thereby causing its death." Willfully to deny a person oxygen which might have prolonged its life has always been a crime. This is, after all, what the Boston stranger did. Dr. Edelin's true "peers" are said to be his fellow abortionists. Why not have the Godfather judged by his fellow Mafiaos?

The jury in Boston (whose religion is unknown and irrelevant except to neo-Nazis) have called to issue that notion that every termination of life done under the rubric of "medical procedure" is not to be tolerated by decent Americans.

EUGENE F. DIAMOND, M.D.  
Chicago

Regarding Dr. Sackler's editorial on the Edelin case, I am surprised at such verbal frothing-at-the-mouth. Dr. Sackler has always seemed like such a calm, cool, deliberate thinker. It's so unlike him. Does he really mean to compare the culpability of food manufacturers in producing coronary disease (a rather far-fetched and tenuous theory at best) with the deliberate actions and inactions of Dr. Edelin? Dr. Edelin, in essence, delivered a premature baby by C-section, and then deliberately registered it to death, by his own admission.

As Dr. Sackler suggests, Dr. Edelin's conviction will probably be overturned—because of technical flaws in his trial—but not because his actions, *per se*, were so noble. He may or may not be guilty of manslaughter, but on the other hand, it ill-behaves so many physicians to make a hero of him, or to publicly applaud his second-trimester "abortion" activities as a prototype of conduct which all physicians should emulate.

Such an attitude is unlikely to rebound to our credit in the future. Out of embarrassment for Dr. Sackler, I will merely pass over his not-too-subtle appeal to religious bigotry, without further elaboration.

As for emotionalism, it surely looks like the shoe is on the other foot this time.

JAMES H. FORD, M.D.  
Lynwood, Calif.

Would sleep with fewer nighttime awakenings benefit your patients with insomnia?

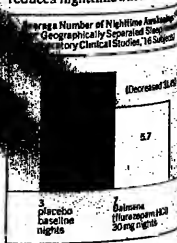
And for those with trouble falling asleep or sleeping long enough...

Dalmane (flurazepam HCl) also delivers excellent results. Clinically proven in sleep research laboratory studies: on average, sleep within 17 minutes that lasts 7 to 8 hours?

Dalmane (flurazepam HCl) is relatively safe, seldom causes morning "hang-over" and is well tolerated. The usual adult dosage is 30 mg h.s., but with elderly and debilitated patients, limit the initial dose to 15 mg to preclude oversedation, dizziness or ataxia. Evaluation of possible risks is advised before prescribing.

Highly predictable results for your patients with trouble staying asleep... can be obtained with Dalmane (flurazepam HCl). As shown below, Dalmane significantly reduces nighttime awakenings.

Average Number of Nighttime Awakenings Geographically Separated Sleep Laboratory Clinical Studies, 16 Subjects



### REFERENCES

1. Kamen L, Williams RL, Smith JR: The sleep laboratory. In: The investigation of sleep and sleep disorders. Scientific exhibit at the 124th annual meeting of the American Psychiatric Association, Washington DC, May 31, 1971.
2. Front 2D In A system for automatically analyzing sleep. Scientific exhibit at the 24th annual Clinical Convention of the American Medical Association, Boston, Nov 20-Dec 2, 1970; and at the 42nd annual scientific meeting of the Aerospace Medical Association, Houston, Apr 26-29, 1971.
3. Nagl GV: Data on file, Medical Department, Hoffmann-La Roche Inc., Nutley NJ.
4. Diment WC: Data on file, Medical Department, Hoffmann-La Roche Inc., Nutley NJ.
5. Data on file, Medical Department, Hoffmann-La Roche Inc., Nutley NJ.

Before prescribing Dalmane (flurazepam HCl), please consult complete product information, a summary of which follows:

- Indications: Effective in all types of insomnia characterized by difficulty in falling asleep, frequent nocturnal awakenings and/or early morning awakenings; in patients with recurring insomnia or chronic medical situations requiring sedation; in patients with anxiety and insomnia.
- Contraindications: Known hypersensitivity to flurazepam HCl.

Depend on highly predictable results with

**Dalmane**  
(flurazepam HCl)

One 30-mg capsule h.s. = usual adult dosage (15 mg may suffice in some patients).  
One 15-mg capsule h.s. = initial dosage for elderly or debilitated patients.

specifically indicated for insomnia

Objectively proved in the sleep research laboratory:  
a sleep with fewer nighttime awakenings  
a sleep within 17 minutes, on average  
a sleep for 7 to 8 hours, on average, with a single h.s. dose.



ROCHE LABORATORIES  
Division of Hoffmann-La Roche Inc.  
Nutley, New Jersey 07110



# We know Librium works. (chlordiazepoxide HCl)

## We're still learning more about how and why.

### Value of continuing animal research

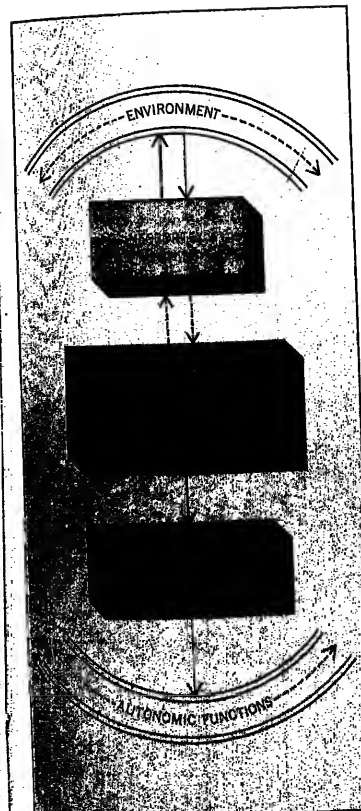
Clinical knowledge of Librium is extensive, yet its mode of action remains under continuing study. Data from animal experiments have been presented here for their intrinsic interest and because such findings often provide direction to new research, both experimental and clinical. However, conclusions from such studies may not always be extrapolated to humans.

### Is the limbic system the "Librium (chlordiazepoxide HCl) system"?

A great deal of experimentation on various animal species suggests that the limbic system is the principal site of action of Librium. Thus, in freely moving cats with electrodes implanted in the brain, Librium 5 mg/kg i.p. slowed electrical activity in the hippocampus, amygdala and septal areas but not in the neocortex which was significantly affected only at higher doses.<sup>1,2</sup> Current investigations on monkeys,<sup>3,4</sup> however, indicate that other subcortical structures may be implicated in the effect of Librium.

Other investigators, through electrophysiologic studies<sup>5</sup> in intact, conscious cats and monkeys, have demonstrated that chlordiazepoxide activates structures involved in the rewarding system—the preoptic area, lateral hypothalamus, septal region and hippocampal formation. At the same time, it appears to inhibit structures implicated in aversive behavior—the thalamic nuclei of the diencephalon and the midbrain reticular formation (MRF).

References:  
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5. Guerrero-Figueroa R, et al: Electrophysiological analysis of the action of four benzodiazepine derivatives on the nervous system, *ibid*, pp. 489-511.



Schematic demonstrating hypothetical pathways of emotional activity and its related expression in laboratory animals.

### Clinical significance of excessive anxiety

Anxiety, when inappropriate and immoderate, may not only have adverse psychological effects but may also cause various somatic disturbances. Reduction of excessive anxiety thus contributes to relief of anxiety-linked emotional and physical disorders.

### Antianxiety action of Librium (chlordiazepoxide HCl)

The dependable action of Librium has been demonstrated in the relief of excessive anxiety and tension occurring alone or in association with functional and organic disorders—usually without adversely affecting performance. Librium is often used concomitantly, when anxiety is a contributing or complicating factor, with certain specific medications of other classes of drugs, e.g., cardiac glycosides, diuretics and antihypertensives.

Adjunctive use of Librium is recommended when counseling, reassurance or other nonpharmacologic measures alone are not considered sufficiently effective. When anxiety has been reduced to manageable levels, therapy with Librium should be discontinued.

**Librium®**  
(chlordiazepoxide HCl)  
5 mg, 10 mg, 25 mg capsules

ROCHE

We're still learning more about it  
to make it more useful to you.

Before prescribing, please consult complete product information, a summary of which follows:

**Indications:** Relief of anxiety and tension occurring alone or accompanying various disease states.

**Contraindications:** Patients with known hypersensitivity to the drug.

**Warnings:** Caution patients about possible combined effects with alcohol and other

CNS depressants. As with all CNS-acting drugs, caution patients against hazardous occupations requiring complete mental alertness (e.g., operating machinery, driving). Though physical and psychological dependence have rarely been reported on recommended doses, one caution in administering to addiction-prone individuals or those who might increase dosage without withdrawal symptoms (including convulsions).

following discontinuation of the drug and similar to those seen with barbiturates, have been reported. Use of any drug in bearing age requires that its potential benefits be weighed against its possible hazards. **Precautions:** In the elderly and debilitated, add in children over six; limit in smallest effective dosage (initially 10 mg or less per day) to produce ataxia or oversedation.

increasing gradually as needed and tolerated. Not recommended in children under six. Though generally not recommended, if combination therapy with other psychotropics seems indicated, carefully consider individual pharmacologic effects, particularly in use of potentiating drugs such as MAO inhibitors and phenothiazines. Observe usual precautions in presence of impaired renal or hepatic function. Par-

adoxical reactions (e.g., excitement, stimulation and acute rage) have been reported in psychiatric patients and hyperactive, aggressive children. Employ usual precautions in treatment of anxiety states with evidence of impending depression; suicidal tendencies may be present and protective measures necessary. Variable effects on blood coagulation have been reported very rarely in patients receiving the drug and

oral anticoagulants; causal relationship has not been established clinically. Adverse Reactions: Drowsiness, ataxia and confusion may occur, especially in the elderly and debilitated. These are reversible in most instances by proper dosage adjustment, but are also occasionally observed at the lower dosage ranges. In a few instances syncope has been reported. Also, encountered are isolated instances of skin

eruptions, edema, minor menstrual irregularities, nausea and constipation, extrapyramidal symptoms, increased and decreased libido—all infrequent and generally controlled with dosage reduction; changes in EEG patterns (low-voltage fast activity) may appear during and after treatment; blood dyscrasias (including agranulocytosis, leukopenia and hepatic dysfunction) have been reported occasionally, making

periodic blood counts and liver function tests advisable during protracted therapy. **Supplied:** Librium® Capsules containing 5 mg, 10 mg or 25 mg chlordiazepoxide HCl. Librium® Tablets containing 5 mg, 10 mg or 25 mg chlordiazepoxide.

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Roche Laboratories  
Division of Hoffmann-La Roche Inc.  
Nutley, New Jersey 07110

## wine talk

By JOHN CHAMBERS  
Author and Consultant to  
Morrell & Company,  
New York Wine Merchants

### Aging Wine

Dr. ARTHUR BIEGANSKI of New York is one of the most ardent wine enthusiasts of my acquaintance. Indeed, at his home it would be almost an insult to ask for Scotch. He has a genius for coming up with surprises, and his latest was a master stroke. Somehow he had managed to find a bottle of ruby port that had lain in a cellar over 30 years. To taste the delicacy and beauty of this comparatively inexpensive generally available wine, given 30 years' aging, was a potent reminder of what you can do for a wine with the capacity in respond to it.

Some wines do not need aging and are best drunk very young. These are light wines—rosés and whites for the most part, and a few of the lightest reds like Beaujolais, Bardolino, or inexpensive Chianti. Only the better white Burgundies and Graves, the Pinot Chardonnays of California, Barsacs and Sauternes, the white Rhones, and the spittles, anisees, and up of Germany need more than a year or two in the bottle, and of these, only the white Rhones, Barsacs, Sauternes, and sweeter German wines can be kept with impunity beyond seven years.

With red wines the problem of aging becomes more complicated. Here it is not only a question of a particular wine, but also the character of a vintage. For example, most of the 1967 red Bordeaux are ready for present drinking, whereas the bigger 1966's are a year or two away. The best rule of thumb is that red Bordeaux from the Médoc require seven years, from St. Emilion and Pomerol six years, and from elsewhere in Bordeaux four years. If the vintage is listed as a "long-lived" one, add a year or two.

### Burgundy Needs Less Time

Red Burgundy is ready sooner. Wines from the Côte de Nuits require six years in a big vintage, whereas most Côte de Beaune are ready in four years. Only the biggest Beaujolais will improve beyond three years. In the Rhone valley the biggest wines require seven years of bottle age in most vintages, but Côtes du Rhone (one of the better buys on the market) need only two to three years. The same holds true of the Loire reds and of the so-called country reds from elsewhere in France.

In Italy, Spain, and Portugal, price is a fair guide to aging requirements. The more expensive wines need six to seven years in the bottle, while two to three years is sufficient for the less expensive. The other red wines of Europe can generally be drunk when marketed.

Most reds from North and South America can also be drunk when purchased, the major exceptions being the better California Cabernet Sauvignons, Petite Sirahs, and Zinfandels, all of which benefit from additional bottle age.

Next Month: Research and Viniculture

## Strike Pact Terms May Have Wide Impact

Continued from page 1

hospital, consisting of an equal number of members from staff and administration, and charged with formulating "appropriate" work schedules" and hearing grievances. Salaries, which now range from \$13,500 for an intern to \$19,200 for a 6th year resident (PGY 7) will rise 8 per cent, with an across-the-board cost of living sum of \$250 added.

### Broad Impact Foreseen

Dr. Robert G. Harmon, president of the Physicians National Housestaff Association, which supported the strike, told MEDICAL TRIBUNE that he thought the point would not be lost on "exploitative" hospital administrators and senior staff everywhere.

"It's a major victory, and it's going to give momentum to National Labor Relations Board negotiations for reasonable hours and working conditions in many hospitals—far example, Los Angeles County Hospital and the District of Columbia Children's Hospital," Dr. Harmon said. He noted that Dr. Malcolm C. Todd, president of the A.M.A., had given his blessing to the strike, a move that surprised some and could not help enhancing the possibility of similar changes, if not strikes, elsewhere.

Dr. Todd's statement said in part that "in important respects, this is a strike for better patient care... When a physician has to work 50 hours straight or 100 hours in a week, it is not only tough on him or her, it is also a threat to the quality of care the patient is receiving."

### Hospitals Reject Implications

A spokesman for the A.M.A. told MEDICAL TRIBUNE that although the Association expected criticism from its membership concerning the Todd statement, little had yet been received. However, officials of the struck hospitals in New York vigorously rejected the implications of the statement, saying that the League of Voluntary Hospitals included some of the finest medical facilities in the world and would never do anything that threatens patient care.



In the first doctors' strike recruited in the U.S., picket lines surrounded some of the most prestigious hospitals in the country, including Mount Sinai, above.

Dr. S. David Pomrinse, director of Mount Sinai Hospital, one of those affected, asserted that long hours are necessary for house officers' training and are not detrimental to interns' and residents' health or to the care they give their patients. He maintained that even on the longest shifts, staffers have time for naps in between cases.

"Our chiefs of service are just as concerned as they are about their health," he said of the interns and residents. "There is adequate time for rest." Until recent years, he said, schedules were even tougher, with house officers being on duty three days and two nights on a regular basis.

Jess Solivan, president of the League, backed Dr. Pomrinse. Regardless of scheduling, he said, "It's expected that when a doctor reaches the point where he's not able to produce or to avail himself of the learning process, he'll say, 'Hey, give me some relief!'"

This just isn't so, contended Dr. Mark Fleischer, a medical intern on the picket line at Brookdale Hospital Medical Center in Brooklyn. "What

are you going to do at 3:00 A.M.? Call your family, who's in the same condition you are and say, 'Hey, give me some relief!'"

An attending physician at one of the struck hospitals confirmed Dr. Fleischer's statements. This physician, who wished to remain anonymous, recalled that when he took his internship at Montefiore Hospital and Medical Center in the Bronx, which was also affected by the strike, some few interns did call for help from their chiefs of service.

### Retaliation Recited

"In most cases they got it," he retorted. "But they always paid for it later. They were branded as weak sisters who couldn't take the strain of being a doctor, and in some cases I know of, they weren't asked back to take their residencies at Montefiore the following year."

And Dr. Don Rubin, a medical intern at Mount Sinai, pointed out that a house officer on a 36- to 48-hour tour

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of duty may actually get less sleep than Dr. Pomrinse did on his 60-hour shifts. "Medicine has become much more complex in recent years," said Dr. Rubin, who took his turn on the picket line at his institution. "There's much more that we can do for patients."

"For instance, when a patient went into cardiac arrest in Dr. Pomrinse's unit, the intern signed the death certificate and went back to bed. Today, he's going to be working with the cardiac emergency team for at least two hours, saving the patient's life."

"And it's the same with peritoneal dialysis, which they didn't have until the early 1960s. If a patient needs dialysis today, he doesn't die. But I'm sitting up all night with him."

### Out-of-Title Work Cited

Dr. Rubin cited the demand for shorter hours to the out-of-title work issue. "A lot of what I do, especially at night, isn't doctor work. Watching that dialysis patient should be done by a nurse, with me on call. And I spend a lot of time wheeling patients around in the hospital or delivering bloods to the lab."

Not all the house officers at League hospitals went out on strike. At Brookdale, for instance, many of the senior medical residents stayed on, while most of their junior colleagues walked the picket line.

"Some of us were angry about that," Dr. Fleischer said. "But in a way, it made things easier on me to know there were doctors in there taking care of the patients."

Most of the striking house officers, led as Dr. Fleischer did, and at many of the struck institutions the house officers made arrangements with the hospital to provide emergency patient care.

"It wasn't unusual" a surgery resident at one such facility said, "to see a picket put down his sign, go into the emergency room to help out with a real crisis, and then come back out and re-

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League of Voluntary Hospitals officials announcing strike settlement. Clockwise from lower left, William A. Abelson, executive director, Jess Solivan, president, Norman Metzger of Mount Sinai Hospital, and Alan Abramson of Montefiore.

sume picketing." Such actions, he noted, are illegal under the 1974 federal law governing hospital strikes.

On the other hand, some residents who did not walk out lent support to the strike from within the hospital. A house officer who remained on the job reported that his colleagues refused to work up patients who were admitted for elective surgery after the strike began.

One major League facility had no strikers at all—the 800-bed New York University Medical Center. House officers there believed that even if attending physicians could take up the slack completely, the Hippocratic Oath forbade a strike.

"This doesn't mean we didn't support the aims of the strike," said one pediatric resident there. "A lot of us did. We just didn't feel that a strike was the right way to do it."

The decision to walk out was a major one for the C.I.R.'s strike committee. "It was forced on us by the

## Tribune Economic Analysis



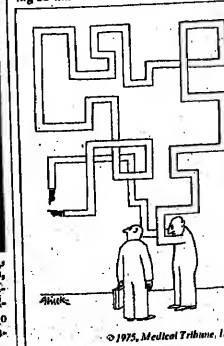
The very magnitude of today's debt burden offers a handle for avoiding a repeat performance of the 1930s' depression that is clearly threatening. Over-indebtedness is the specific abuse responsible for hyper-inflation. Booms invite an overload of debt, which accentuates busts.

The sound way to undo today's damage, and to avoid still more, is to lighten the debt load by trading on the troublesome fact that people on payrolls are struggling with every bit as eroded a debt overload as outlays to meet payrolls. The banks are at least as anxious over the consequences of their over-lending as their debtors are over the consequences of their over-borrowing.

### A 3-Way Compromise

The wobbling companies, their nations banks, and the petrified people on their payrolls would all be ahead if they worked a three-way compromise. Assume that the management in trouble could show both the banks on its back and the people on its payroll how much difference a reasonable cut would make. And that management demonstrated its good faith by practicing austerity on expense accounts and taking an appropriate cut itself. All three partners in the debt squeeze would be ahead if management "borrowed" the pay cut from labor instead of just taking it.

Issuing company notes to everyone on the payroll in order to cover the cut agreed upon would kill three birds with one stone. Management would cut costs. People now worrying that each paycheck might be the last would get a new asset with a fighting chance to keep the money coming. The banks would wind up with a better-fixed business borrower, plus a whole new group of family circle customers for the consumer installment loans they are pushing so hard.



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Delegates of Committee of Interns and Residents take a straw vote on contract offer by the League of Voluntary Hospitals during negotiations.

C.I.R. Photo

# "Let me tell you about the medicine I'm going to prescribe."

## TALKING OVER VALIUM®(diazepam) THERAPY WITH YOUR ANXIOUS PATIENT.



And it's also good for him to realize that he will be taking Valium only as long as he needs it.

Your expressed confidence in the medication prescribed, and the positive atmosphere in which therapy is given and accepted, work to the patient's advantage.

A patient often benefits by a greater understanding of his treatment program. You may find it helpful to make your patient aware that the purpose of therapy with Valium is to help reduce discomforting and disabling symptoms of excessive psychic tension and anxiety. It is beneficial for him to understand that much of his tension and anxiety can be relieved by your reassurance and counseling, and that these measures can do more than anything else to help him cope with his basic problems. The patient is reassured in knowing he can expect his medication to help him avoid feeling overwhelmed by his symptoms.

Selection of a dosage regimen is an important consideration when Valium (diazepam) is prescribed, and dosage should be individualized to achieve maximum beneficial effect. If the patient understands clearly when and how much to take, and if he knows why it's to his benefit to follow the regimen closely, the chances are better that he will take the medication precisely as directed. That should help avoid missed doses and discourage taking too much or too little medication — all of which can have an undesirable effect on the management of the patient's condition.

*"It's important that you follow my directions closely."*

*"I'll see you again the week after next and we'll see how you're making out."*

Your patient is often likely to feel reassured when you talk about seeing him again to check his progress. A planned visit evidences your continued interest and affords the patient an opportunity to report improvement he has made and to relate whatever continuing or additional difficulties he may be experiencing. It's also a chance for him to describe his response to therapy with Valium.

During follow-up visits, as your patient talks about his medication and about its effects on his symptoms, he will provide the kind of information that will be of great help in evaluating total therapy, adjusting the dosage of Valium, or discontinuing the medication entirely if that seems indicated.

## Valium® (diazepam)

2-mg, 5-mg, 10-mg scored tablets  
for individualized treatment of psychic tension



Please see the following page for a summary of product information.





# Valium® (diazepam)

2-mg, 5-mg, 10-mg scored tablets

**Prompt, effective action.** Valium (diazepam) works rapidly to relieve pronounced psychic tension in patients overreacting to stress and in psychoneurotic patients.

Before prescribing, please consult complete product information, a summary of which follows:

**Indications:** Tension and anxiety states; somatic complaints which are concomitants of emotional factors; psychoneurotic states manifested by tension, anxiety, apprehension, fatigue, depressive symptoms or agitation; symptomatic relief of acute agitation, tremor, delirium tremens and hallucinosis due to acute alcohol withdrawal; adjunctively in skeletal muscle spasm due to reflex spasm to local pathology; spasticity caused by upper motor neuron disorders, athetosis, stiff-man syndrome; convulsive disorders (not for sole therapy).

**Contraindicated:** Known hypersensitivity to the drug. Children under 6 months of age. Acute narrow angle glaucoma; may be used in patients with open angle glaucoma who are receiving appropriate therapy.

**Warnings:** Not of value in psychotic patients. Caution against hazardous occupations requiring complete mental alertness. When used adjunctively in convulsive disorders, possibility of increase in frequency and/or severity of grand mal seizures may require increased dosage of standard anticonvulsant medication; abrupt withdrawal may be associated with temporary increase in frequency and/or severity of seizures. Advise against simultaneous ingestion of alcohol and other CNS depressants. Withdrawal symptoms (similar to those with barbiturates and alcohol) have occurred following abrupt discontinuance (convulsions, tremor, abdominal and muscle cramps, vomiting and sweating). Keep addiction-prone individuals under careful surveillance because of their predisposition to habituation and dependence. In pregnancy, lactation or women of childbearing age, weigh potential benefit against possible hazard.

**Precautions:** If combined with other psychotropics or anticonvulsants, consider carefully pharmacology of agents employed; drugs such as phenothiazines, narcotics, barbiturates, MAO inhibitors and other anti-

Wide margin of safety. Valium is generally well tolerated and in usual dosages rarely produces significant adverse reactions. (See prescribing information below.)

**Dosage flexibility.** Scored Valium 2-, 5-, and 10-mg tablets give you dosage flexibility no tranquilizer capsule can match.

depressants may potentiate its action. Usual precautions indicated in patients severely depressed, or with latent depression, or with suicidal tendencies. Observe usual precautions in impaired renal or hepatic function. Limit dosage to smallest effective amount in elderly and debilitated to preclude a taxia or oversedation.

**Side Effects:** Drowsiness, confusion, diplopia, hypotension, changes in libido, nausea, fatigue, depression, dysarthria, jaundice, skin rash, ataxia, constipation, headache, incontinence, changes in salivation, slurred speech, tremor, vertigo, urinary retention, blurred vision. Paradoxical reactions such as acute hyperexcited states, anxiety, hallucinations, increased muscle spasticity, insomnia, rage, sleep disturbances, stimulation have been reported; should these occur, discontinue drug. Isolated reports of neutropenia, jaundice; periodic blood counts and liver function tests advisable during long-term therapy.

**Dosage:** Individualize for maximum beneficial effect. **Adults:** Tension, anxiety and psychoneurotic states, 2 to 10 mg b.i.d. to q.i.d.; alcoholism, 10 mg t.i.d. or q.i.d. in first 24 hours, then 5 mg t.i.d. or q.i.d. as needed; adjunctively in skeletal muscle spasm, 2 to 10 mg t.i.d. or q.i.d.; adjunctively in convulsive disorders, 2 to 10 mg b.i.d. to q.i.d. **Geriatric or debilitated patients:** 2 to 2½ mg, 1 or 2 times daily initially, increasing as needed and tolerated. (See Precautions.) **Children:** 1 to 2½ mg t.i.d. or q.i.d. initially, increasing as needed and tolerated (not for use under 6 months).

**Supplied:** Valium® (diazepam) Tablets, 2 mg, 5 mg and 10 mg—bottles of 100 and 500; Tel-B-Dose® packages of 100, available in trays of 4 reverse-numbered boxes of 25, and in boxes containing 10 strips of 10; Prescription Paks of 50, available singly and in trays of 10.

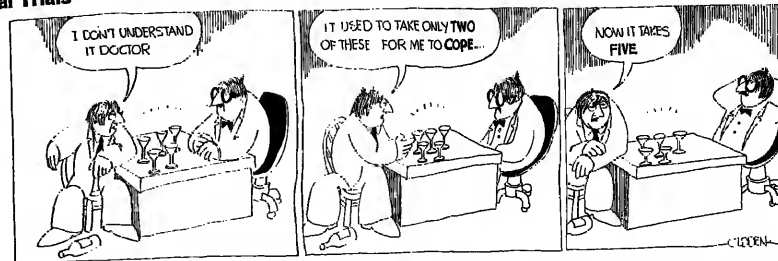


Roche Laboratories  
Division of Hoffmann-La Roche Inc.  
Nutley, New Jersey 07110

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MEDICAL TRIBUNE

## Clinical Trials



## TRIBUNE SPORTS REPORT

### Hang Gliding Said to Point Up Need for 'Action Priorities'

Medical Tribune Report

SAN FRANCISCO—Dr. Arthur E. Ellison of Williamstown, Mass., cited the fast-growing sport of hang gliding as an example of the need to establish "action priorities" in athletic medicine through coordination of research efforts.

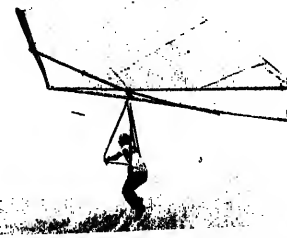
He told a meeting of the American Orthopaedic Society for Sports Medicine here that participants in this dangerous pastime have increased from 200 to 11,000 since 1972, with gliding kites now being sold at a rate of about 1,000 a month.

#### A Death a Month in California

While the exact injury rate is not known, he said, Rancho Los Amigos in Los Angeles has six paraplegic patients who are victims of hang gliding accidents, and California alone averages one fatality a month from this sport.

Most of the injuries are to the extremities, he said. As some of the factors in the accidents, Dr. Ellison cited the weather,

the terrain, pilot error, and kite failure. A thorough survey, he said, might suggest that a special action program is required, including: modification of equipment to provide such safety devices as a parachute or an ejection suit; special padding, helmets, gloves, or boots; elimination of flying over dangerous terrain; licensing; a ban on unsound kites; or, if the toll is found to be too high, outlawing of hang gliding altogether.



### Calif. Acupuncture Unit With Few MDs Likely

Continued from page 9

puncture which is lobbying to change the Moscone bill.

"There is room for traditional acupuncturists," Dr. Wong said in an interview with MEDICAL TRIBUNE, "but they should be required to show expertise in basic science. By the same token, we also think that M.D.s should not be given carte blanche, but ought to be required to take 100 to 150 hours of acupuncture training, as they must do now in New York."

#### Alternative Makeup Proposed

"As for the makeup of the Advisory Board, our alternative suggestion is a 10-member Board with the following distribution: one member from the State Board of Medical Examiners; four physician-acupuncturists; one dentist-acupuncturist; one non-medical, academic, research-oriented Ph.D. with at least five years experience with acupuncture; and three traditional acupuncturists trained in Japan, China, or Korea; with at least ten years experience, and demonstrated knowledge of western concepts of anatomy, physiology, etc."

"Our main motive," he added, "is to see that the public is fully protected,

and acupuncture doesn't go 'down the pipes' as quackery."

Sources in Gov. Brown's office told MEDICAL TRIBUNE that he is waiting to study the final version of the Moscone bill before deciding whether to sign it; they said the Assembly often amends or adds to bills received from the Senate.

Neighboring Nevada, in 1973, was the first state to legalize the practice of acupuncture by non-physicians without medical supervision. In the rest of the country, there is a patchwork of regulations, often stipulating that acupunc-

ture can only be performed by M.D.'s for research purposes.

The A.M.A. has not adopted an official policy on acupuncture. However, the August, 1974 statement issued by the August, 1974 statement issued by the A.M.A. delegation on its return from the Peoples' Republic of China said that "acupuncture analgesia merits its controlled experimental study," while warning that "acupuncture therapy should be regarded as the practice of medicine in an experimental phase, permissible only in qualified investigative settings."

### Gonorrhea in Women Declared to Be Often Symptomatic

Medical Tribune World Service

GENOVA—Gonorrhea in women cannot be regarded as commonly nonsymptomatic, a United States physician stated here at a World Health Organization-sponsored meeting on health education in the control of sexually transmitted diseases.

Estimates that up to 60 per cent of infected women, and 10-20 per cent of men, are without symptoms, are largely based on the experience of physicians working in VD clinics, said Dr. King K. Holmes, of University of Washington, Seattle.

Such views, he said, are not corroborated by probability sampling, case control, or cohort studies, or from the syndrome-oriented experience of certain subspecialties, such as rheumatology and gynecology.

#### Asymptomatic in 80%

"About 80 per cent of women seen in the University of Washington specialty clinics and emergency room have sought treatment because of acute symptoms," Dr. Holmes said.

As manifestations in women that are suggestive of, or compatible with, gon-

by G.

## IMMATERIA MEDICA

### Minnesota Medicine's Mascot

The new editor-in-chief of *Minnesota Medicine*, Dr. Richard L. Reeves, has introduced a mascot into his columns. Why? "Because I have one in mind, that's why," says Editor Reeves. "His name is minny."

That brought us up short. Unless, we figured, for *Minnesota*. Editor Reeves says minny is "a literary cockroach who composes free verse by hurling himself head downward against the typewriter keys."

Like Don Marquis' archbald, of *Archbald and McWhorter* fame, from whom minny is descended, he can't manage capital letters or punctuation on the typewriter. "Minny is bold, disrespectful, fun-loving, contemptuous of detail, and hungry for the literary life," says Editor Reeves, who in his March issue published minny's first poem:

my minnesota medicine editor  
i accept the position  
because mascots bring luck  
and you will need plenty

Long live minny the mascot of *Minnesota Medicine*! Who knows? This mascot business may be as contagious as measles. We could have minny for *Virginia Med. M.*, for *Florida M.A.*, for *Pennsylvania Med.*, for *Texas M.*, for *Missouri Med.*, and missy for *Mississippi Med. Ass.* But now that they are teaching chimps to talk and typewrite, nobody says all mascots have to be cockroaches. In fact, we know some who are just cute nurses.

While, currently, 10 to 20 per cent of male patients at VD clinics have no symptoms, this figure also bears no relationship to the true proportion of new cases of this kind, Dr. Holmes asserted.

"In an unpublished cohort study, we have found this proportion to be only 3 per cent," he reported.